CASE REPORT

Acute Pancreatitis Complicated by Jejunal Hematoma in a Patient on Anti-Coagulants and Anti-Platelets

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ABSTRACT

Context Pancreatitis can be associated with significant complications. Bowel hematoma is a rare complication and the second part of the duodenum is the most commonly affected site. Hematomas affecting other parts of the bowel are extremely rare. Case report A 53-year-old female with a history of atrial fibrillation and ischemic heart disease on anticoagulants and aspirin presented with abdominal pain of a few days duration which had worsened prior to presentation. This was associated with abdominal distension, vomiting and melena. Laboratory investigations showed elevated serum amylase, coagulopathy and severe anemia. Computed tomography imaging showed a jejunal hematoma and pancreatitis with peripancreatic inflammation. She responded to conservative treatment in addition to correction of the coagulopathy and a blood transfusion. Her symptoms were resolved within a few days and a repeat computed tomography scan two months later showed complete resolution of the hematoma and the pancreatitis. Conclusion Our case demonstrates a rare complication of acute pancreatitis in a patient with risk factors.

INTRODUCTION

Acute pancreatitis can be associated with significant complications and these can be categorized into either early or late and local or systemic. [1, 2] Significant local complications include necrosis, pseudocysts and abscess whereas significant systemic complications include shock, respiratory failure, renal failure, metabolic derangements and disseminated intravascular coagulopathy. When inadequately treated, these complications are associated with a high mortality rate. Hemorrhagic complications are less common and these include gastrointestinal bleeding which can be immediate or delayed, a spontaneous splenic hematoma and a bowel hematoma [3, 4, 5]. In the case of a bowel hematoma, the duodenum is the most commonly affected site. We report a rare case of a jejunal hematoma secondary to acute gallstone pancreatitis in a patient who was on anti-coagulants and anti-platelets for atrial fibrillation and ischemic heart disease.

CASE REPORT

A 53-year-old female presented to the Emergency Room with a two-day history of hematuria and melena, and a one day history of worsening central abdominal pain, distension and vomiting of bilious fluid. She had initially experienced epigastric and right upper quadrant pain a few days before her latest symptoms. This pain resolved without treatment and she mentioned that the subsequent pain was different in character from the initial pain. Her past medical history included thyrotoxicosis under treatment, hypertension, ischemic heart disease and atrial fibrillation. Her medications consisted of carbimazole, enteric-coated aspirin, omeprazole, atenolol and warfarin. She denied using any other medications in the weeks before this current illness. There was definitely no history of abdominal trauma or any flu-like illness before the onset of the symptoms. On examination, she looked pale and had tachycardia of 140 beats per minute (irregular) but was maintaining her blood pressure at 130/70 mmHg. Abdominal examination showed a distended abdomen which was not tender and not guarded. Laboratory investigations showed severe anemia (hemoglobin 5.2 g/dL, reference range: 12-16 g/dL; leukocytosis 15.0 x10^9/L, reference range: 4.0-11.0 x10^9/L), coagulopathy higher than the designated therapeutic target (international normalized ratio, INR 4.7) and mild renal impairment (serum creatinine 171 μmol/L, reference range: 35-88 μmol/L). Serum calcium and lipid profiles were normal. When she had
attended her routine anti-coagulation clinic appointment the previous day, her INR was 6.5 and the warfarin was stopped. Apart from this, her anticoagulation control had previously been satisfactory. At that point, her abdominal symptoms were still bearable. Abdominal radiography showed small dilated fluid-filled bowel loops suggesting bowel obstruction. Her urine was clear but urinalysis detected red blood cells.

She was immediately referred to the surgical service and an urgent computed tomography (CT) scan was carried out. This showed a thickened jejunal loop consistent with a hematoma, a dilated fluid-filled proximal duodenum and stomach, a solitary gallstone and a swollen pancreatic head and body with peripancreatic fat streaking, consistent with acute pancreatitis (Figure 1). The jejunal hematoma was located in the proximal jejunum and extended for approximately 10 cm. Close inspection of the pancreatic and common bile duct did not reveal any abnormalities, such as dilatation, stones or pancreas divisum. There were no abnormalities, such as calculus, seen in the urinary system to account for the hematuria. Laboratory investigations revealed an elevated serum amylase level of 1,962 IU/L (reference range: 0-120 IU/L) and normal liver function with the exception of elevated serum bilirubin of 78 μmol/L (reference range: 0-34 μmol/L). A review of her previous liver function tests carried out when she had previously complained of intermittent mild epigastric pain had shown intermittent cholestatic profiles which had resolved spontaneously. Unfortunately, she was not evaluated with an ultrasound scan of the abdomen. Given the history of abdominal pain characteristic of biliary colic which had preceded the abdominal distension, a diagnosis of acute gallstone pancreatitis complicated by a jejunal hematoma was made. This was despite the minimally abnormal liver profiles on admission.

As the patient was mildly symptomatic and stable, she was referred to the gastroenterology service and was admitted to the general medical ward. She was kept nil orally, started on intravenous fluids and omeprazole, and given blood and fresh frozen plasma transfusions to correct the anemia and the coagulopathy. Her condition remained stable and her symptoms improved over the next few days. Her abdominal distension and vomiting regressed and she was allowed oral feeding. Echocardiography showed a normal left atrium without any evidence of clots. She was discharged seven days after admission but was not started on her aspirin and warfarin. Repeat urinalysis before discharge showed resolution of the microscopic hematuria. A repeat CT scan two months later showed that the hematoma and the other abnormalities seen on the initial CT scan had completely regressed (Figure 2). She has begun her anti-coagulants again with close monitoring and has remained well on follow-up without further abdominal pain or hematuria.

**DISCUSSION**

Bowel hematoma can be categorized as either spontaneous or traumatic. A traumatic hematoma is by far the most common and is often associated with blunt trauma injuries or found after endoscopy with biopsies [6, 7, 8]. However, in the latter, risk factors, such as
hematological disorders, liver cirrhosis or anti-platelet agents, are usually present. Spontaneous or non-traumatic hematomas are most commonly associated with over-anticoagulation, hematological disorders, vascular disorders or malignancies [9, 10, 11, 12, 13, 14]. Pancreatitis is a recognized but rare cause of bowel hematoma [5, 15, 16]. The duodenum is the most commonly affected site in acute pancreatitis due to its close proximity to the pancreas. The exact mechanism underlying the formation of a bowel hematoma in pancreatitis is unknown and can vary depending on the underlying etiology. Two hypotheses have been postulated by van Spreeuwel et al. [17]. The first hypothesis involves the presence of ectopic pancreatic tissue within the bowel wall. Similar to normal pancreatic tissue, an ectopic pancreas can develop acute inflammation, and the subsequent necrosis can lead to hematoma formation. However, ectopic pancreatic tissue is rare and is most commonly located in the antrum of the stomach. The second mechanism postulated suggested that the leakage of the pancreatic enzymes in pancreatitis can result in tissue damage leading to local necrosis and hematoma formation. However, these mechanisms have only been postulated for a duodenal hematoma in association with pancreatitis. Whether similar events can lead to a bowel hematoma in another part of the bowel is not known. In our patient, there was prominent peripancreatic inflammation in close proximity to the proximal portion of the jejunal hematoma. This suggests that the pancreatitis contributed to the jejunal hematoma. The use of anti-platelets and anti-coagulants, which were in excess of the correct therapeutic dosage at that time, could have contributed greatly to the underlying pathogenesis in our patient.

The management of a jejunal hematoma is similar to the management of a bowel hematoma in another location. In some cases, surgery has been performed mainly because the diagnoses were not known pre-operatively [9]. As most hematomas will resolve with conservative and supportive treatment, non-surgical interventions are currently recommended [9]. Keeping patient nil orally, intravenous supplements and corrections of coagulopathy and anemia are the mainstays of treatment. In most cases, the hematoma, depending on its size, will resolve in a few days. Our patient had complete obstruction of the jejunum and started to feel better within a few days of conservative treatment even before the hematoma had regressed significantly. This is not surprising as even minor resolution of a hematoma allows re-establishment of lumen patency in the patient, thus allowing symptom improvement. Interestingly, apart from being caused by a bowel hematoma, acute pancreatitis can be caused by a duodenal hematoma due to compression of the pancreatic duct [18, 19]. Therefore, in certain cases, it may be difficult to distinguish which of these events came first. In our case, given the history of colicky epigastric and right upper quadrant pain preceding the bowel distension by a few days, we are certain that the pancreatitis was the initial event leading to the hemorrhagic complication. Furthermore, the hematoma was localized in the jejunum, too far away to cause pancreatic duct obstruction.

In conclusion, our case points out that a hematoma can occur in any part of the gastrointestinal tract in close proximity to the pancreas, especially in patients with risk factors, such as anticoagulant and anti-platelet therapy. Management includes standard treatment for bowel obstruction and correction of the coagulopathy and anemia.

Conflicts of interest The authors have no potential conflict of interest

References


